#### M.Sc.

#### 2014

# 2nd Semester Examination

# PAPER-BLM-201 (UNIT-10)

Full Marks: 40

BIOMEDICAL LABORATORY SCIENCE AND MANAGEMENT

Time: 2 Hours

The figures in the right-hand margin indicate full marks.

Candidates are required to give their answers in their own words as far as practicable.

Illustrate the answers wherever necessary.

Answer all questions.

## (Module - 1)

- 1. Answer any five questions of the following:  $5 \times 1$ 
  - (a) Which one of the statement is true about haemoglobin:
    - (i) One Molecule contains 1 atom of iron;
    - (ii) It has a low level in blood at birth compared with adult life;
    - (iii) It's type is almost same in children and adults.

(D)	total body iron:				
		trandferhin;			
	(ii)	Neutrophil;			
	(iii)	Maenophage.			
(c)		nich of the following is not true about sideroblasti			
	(i)	Ring sideroblast in bone marrow;			
	(ii)	It is most frequently caused by myelodysplasia			
	(iii)	It may be caused by folate deficiency.			
(d)	Which one of the following is used to monito transfusion iron overload:				
	(i)	Serum ferritin;			
	(ii)	Lung function;			
	(iii)	Haemoglobin conten.			
(e)	Complete saturation of oxygenation is found in :				
	(i)	T-form;			
	(ii)	R-form;			
	(iii)	ньг.			

ĺ	ſ	Henostasis	correlates	with	:
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- (i) Thrombocytes;
- (ii) Lymphocytes;
- (iii) Normocytes.

### (g) Truth that Sickle cell anaemia is:

- (i) Glutamic acid replaced by valine;
- (ii) at the  $\beta_6$  perition valine is replaced by glutamine;
- (iii) West Bengal is a high prevalence state.
- (h) Sodium fluoride is used as:
  - (i) anticoagulant;
  - (ii) preservative that prevents red cell lysis;
  - (iii) to prevent glycolysis in red cell.
- 2. (a) Describe the merits of HbF over HbA with justification.
  - (b) Describe the primary, secondary and tertiary structure of globin chains in haemoglobin. 4+4

Or

Describe the significance of the following types of red cells with a clean diagram: 4×2

- (a) Anisocytes;
- (b) Poikilocytes;
- (c) Basophilic strippling;
- (d) Target cells.

- 3. (a) What is crises?
  - (b) State the mechanism of destruction of RBC in sickle cell anaemic patient;
  - (c) What is the significance of red cell indices determination? 2+3+2

Or

- (a) State the role of hepcidin in the regulation of erythropoiesis.
- (b) What is multiple myeloma?
- (c) Classify the types of Beta thalassemia focusing its haemoglobin variants and genetic feature.

2+1+(2+2)

## (Module - 2)

- **4.** Answer any five questions from the following:  $5 \times 1$ 
  - (a) What is haemochromatosis?
  - (b) Write the full form of RDW.
  - (c) What is favism?
  - (d) Why are males mostly affected by G-6-PD deficies than female?
  - (e) What is the normal value MCV?
  - (f) What is the size of the aperture in a coulter-counter chamber?

- (g) What do you mean by microcytic hypochromic anaemia?
- (h) What is meant by thromocytopenia?
- 5. (a) Describe the principle of light scattering system with special reference to FACS for the measurement of different types of blood cells.
  - (b) What is the function of floating calibrator in Coultercountering chamber? 6+2

Or

- (a) How do you interpret your result of paper chectrophoresis in the detection of haemoglobin variants.
- (b) How HBA<sub>1</sub>C is formed?
- (c) Mention the principle of HbA<sub>1</sub>C detection by HPLC.
  3+3+2
- 6. (a) Describe why osmotic fragility test is performed.
  - (b) How do you interpret your test result of osmotic fragility?
  - (c) Mention the significance of reticulocyte count.

3+2+2

Or

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Write short notes on:

- (a) Sideroblastic anaemia;
- (b) Total iron binding capacity;
- (c) Hemophilia.

3+2+2